**ORIGINAL** 

Application Based on

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Inventors: Alexandra DeLauro Bermel and Lori Jeanne Shaw-Klein

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# INK JET RECORDING ELEMENT

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## INK JET RECORDING ELEMENT

## CROSS REFERENCE TO RELATED APPLICATIONS

Reference is made to commonly assigned, co-pending U.S. Patent

5	Applications:			
	Serial Number	by Bermel et al., (Docket 81820) filed of even date		
	herewith entitled "Ink Jet Recording Element";			
	Serial Number	_ by Bermel et al., (Docket 82109) filed of even date		
	herewith entitled "Ink Jet I	Recording Element";		
10	Serial Number	_ by Bermel et al., (Docket 82110) filed of even date		
	herewith entitled "Ink Jet Recording Element";			
	Serial Number	_ by Bermel et al., (Docket 82133) filed of even date		
	herewith entitled "Ink Jet I	Printing Method";		
	Serial Number	_ by Bermel et al., (Docket 82134) filed of even date		
15	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Bermel et al., (Docket 82138) filed of even date		
	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Bermel et al., (Docket 82139) filed of even date		
	herewith entitled "Ink Jet l	Printing Method";		
20	Serial Number	_ by Lawrence et al., (Docket 81815) filed of even date		
	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Lawrence et al., (Docket 81817) filed of even date		
	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Lawrence et al., (Docket 81818) filed of even date		
25	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Lawrence et al., (Docket 81821) filed of even date		
	herewith entitled "Ink Jet Printing Method";			
	Serial Number	_ by Lawrence et al., (Docket 81893) filed of even date		
	herewith entitled "Ink Jet I	Printing Method";		

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Serial Number	by Lawrence et al., (Docket 81894) filed of even date				
herewith entitled "Ink Jet Printing Method"; and					
Serial Number	by Lawrence et al., (Docket 81983) filed of even date				
herewith entitled "Ink Jet Printing Method".					

#### FIELD OF THE INVENTION

The present invention relates to a porous ink jet recording element.

#### BACKGROUND OF THE INVENTION

In a typical ink jet recording or printing system, ink droplets are ejected from a nozzle at high speed towards a recording element or medium to produce an image on the medium. The ink droplets, or recording liquid, generally comprise a recording agent, such as a dye or pigment, and a large amount of solvent. The solvent, or carrier liquid, typically is made up of water and an organic material such as a monohydric alcohol, a polyhydric alcohol or mixtures thereof.

An ink jet recording element typically comprises a support having on at least one surface thereof an ink-receiving or image-receiving layer, and includes those intended for reflection viewing, which have an opaque support, and those intended for viewing by transmitted light, which have a transparent support.

An important characteristic of ink jet recording elements is their need to dry quickly after printing. To this end, porous recording elements have been developed which provide nearly instantaneous drying as long as they have sufficient thickness and pore volume to effectively contain the liquid ink. For example, a porous recording element can be manufactured by cast coating, in which a particulate-containing coating is applied to a support and is dried in contact with a polished smooth surface.

When a porous recording element is manufactured, it is difficult to co-optimize the image-receiving layer surface appearance and ink drying times. Good image-receiving layer surface appearance is obtained when it is virtually crack-free. A crack-free surface appearance can be obtained merely by adding more binder to the image-receiving layer. However, adding more binder increases dry time since the binder fills the pores in the image-receiving layer. Therefore, it is difficult to obtain an image-receiving layer which has a crack-free surface yet is fast-drying.

Another problem encountered with a recording element is ink coalescence which occurs when adjacent ink dots coalesce which leads to nonuniform density.

U.S. Patent 6,037,050 and EP 888,904 relate to an ink jet recording element wherein an ink absorption layer comprises inorganic particles such as silica and a poly(vinyl alcohol) binder that is crosslinked with a hardener.

However, there is no disclosure in these references that the crosslinker should be present in an amount greater than 10%, based on the binder.

It is an object of this invention to provide a porous ink jet recording element that exhibits good overall appearance without cracking and has an excellent dry time and reduced ink coalescence.

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#### SUMMARY OF THE INVENTION

These and other objects are achieved in accordance with the invention which comprises an ink jet recording element comprising a support having thereon a porous image-receiving layer comprising particles, a poly(vinyl alcohol) binder and a crosslinking agent, the particles having a primary particle size of from about 7 to about 40 nm in diameter which may be aggregated up to about 300 nm, and the crosslinking agent being present in an amount of at least about 20 weight % of the poly(vinyl alcohol) binder.

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By use of the invention, a porous ink jet recording element is obtained that exhibits good overall appearance without cracking and has an excellent dry time and reduced ink coalescence.

#### DETAILED DESCRIPTION OF THE INVENTION

Examples of particles useful in the invention include alumina, boehmite, clay, calcium carbonate, titanium dioxide, calcined clay, aluminosilicates, silica, barium sulfate, or polymeric beads. The particles may be porous or nonporous. In a preferred embodiment of the invention, the particles are metallic oxides, preferably fumed. While many types of inorganic and organic particles are manufactured by various methods and commercially available for an image-receiving layer, porosity of the ink-receiving layer is necessary in order to obtain very fast ink drying. The pores formed between the particles must be sufficiently large and interconnected so that the printing ink passes quickly through the layer and away from the outer surface to give the impression of fast drying. At the same time, the particles must be arranged in such a way so that the pores formed between them are sufficiently small that they do not scatter visible light.

The particles may be in the form of primary particles, or in the form of secondary aggregated particles. The aggregates are comprised of smaller primary particles about 7 to about 40 nm in diameter, and being aggregated up to about 300 nm in diameter. The pores in a dried coating of such aggregates fall within the range necessary to ensure low optical scatter yet sufficient ink solvent uptake.

Any fumed metallic oxide particles may be used in the invention. Examples of such particles include fumed alumina, silica, titania, cationic silica, antimony(III) oxide, chromium(III) oxide, iron(III) oxide, germanium(IV) oxide, vanadium(V) oxide, or tungsten(VI) oxide. Preferred examples of fumed metallic oxides which may be used in the invention include silica and alumina fumed

oxides. Fumed oxides are available in dry form or as dispersions of the aggregates mentioned above.

The process for fuming metallic oxides is well known in the art. For example, reference may be made to Technical Bulletin Pigments, no. 56, Highly Dispersed Metallic Oxides Produced by the AEROSIL ® Process, by Degussa AG., 1995.

Any poly(vinyl alcohol) may be used in the invention. In a preferred embodiment, the poly(vinyl alcohol) has an average viscosity greater than about 20 cp when employed in a 4% aqueous solids solution at 20°C. Specific examples of such poly(vinyl alcohols) which may be used in the

Table 1

	Poly(vinyl alcohol)	Average Viscosity @ 4% (cp)
PVA-A	Gohsenol® GH-17	30 <sup>1</sup>
PVA-B	Gohsenol® GH-23	521
PVA-C	Gohsenol® N300	27.5 <sup>1</sup>
PVA-D	Elvanol® 52-22	23.5 <sup>2</sup>

<sup>&</sup>lt;sup>1</sup>Trade publication, Nippon Gohsei Co., Ltd.

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invention include the following:

The amount of poly(vinyl alcohol) binder used should be sufficient to impart cohesive strength to the image-receiving layer, but as small as possible so that the interconnected pore structure formed by the aggregates is not filled in by the binder. In a preferred embodiment of the invention, the weight ratio of the binder to the particles is from about 1:20 to about 1:5.

The image-receiving layer may also contain a mordant. Examples of mordants which may be used include water-soluble cationic polymers, metal salts, water-insoluble cationic polymeric particles in the form of a latex, water dispersible polymer, beads, or core/shell particles wherein the core is organic or inorganic and the shell in either case is a cationic polymer. Such particles can be products of addition or condensation polymerization, or a combination of both.

<sup>&</sup>lt;sup>2</sup>Trade publication, DuPont Corp.

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Examples of crosslinkers which may be used in the invention include carbodiimides, polyfunctional aziridines, aldehydes, isocyanates, epoxides, polyvalent metal cations, acetals, ketals, etc. In a preferred embodiment of the invention, the crosslinker is an aldehyde, an acetal or a ketal. In a more preferred embodiment, the crosslinker is 2,3-dihydroxy-1,4-dioxane.

As noted above, the amount of crosslinking agent is present in an amount of at least about 20 weight % of the poly(vinyl alcohol) binder. This amount is far beyond a typical amount of crosslinking agent for poly(vinyl alcohol). For example, in Paper Coating Additives, Robert J. Kane, TAPPI PRESS, Atlanta Ga., 1995, page 96, it is disclosed that a typical aldehyde crosslinker, glyoxal, is used at about 10% by weight relative to the poly(vinyl alcohol). In a preferred embodiment of the invention, the crosslinking agent is present in an amount of at least about 40 weight %, more preferably about 50 weight %, of the poly(vinyl alcohol) binder.

Since the image-receiving layer is a porous layer comprising particles, the void volume must be sufficient to absorb all of the printing ink. For example, if a porous layer has 60 volume % open pores, in order to instantly absorb 32 cc/m² of ink, it must have a physical thickness of at least about 54  $\mu$ m.

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The support for the ink jet recording element used in the invention can be any of those usually used for ink jet receivers, such as resincoated paper, paper, polyesters, or microporous materials such as polyethylene polymer-containing material sold by PPG Industries, Inc., Pittsburgh,

polymer-containing material sold by PPG Industries, Inc., Pittsburgh, Pennsylvania under the trade name of Teslin ®, Tyvek ® synthetic paper (DuPont Corp.), and OPPalyte® films (Mobil Chemical Co.) and other composite films listed in U.S. Patent 5,244,861. Opaque supports include plain paper, coated paper, synthetic paper, photographic paper support, melt-extrusion-coated paper, and laminated paper, such as biaxially oriented support laminates. Biaxially oriented support laminates are described in U.S. Patents 5,853,965; 5,866,282; 5,874,205; 5,888,643; 5,888,681; 5,888,683; and 5,888,714, the disclosures of which are hereby incorporated by reference. These biaxially oriented supports include a paper base and a biaxially oriented polyolefin sheet, typically polypropylene, laminated to one or both sides of the paper base. Transparent supports include glass, cellulose derivatives, e.g., a cellulose ester, cellulose triacetate, cellulose diacetate, cellulose acetate propionate, cellulose acetate butyrate; polyesters, such as poly(ethylene terephthalate), poly(ethylene naphthalate), poly(1,4-cyclohexanedimethylene terephthalate), poly(butylene terephthalate), and copolymers thereof; polyimides; polyamides; polycarbonates; polystyrene; polyolefins, such as polyethylene or polypropylene; polysulfones; polyacrylates; polyetherimides; and mixtures thereof. The papers listed above include a broad range of papers, from high end papers, such as photographic paper to low end papers, such as newsprint. In a preferred embodiment,

The support used in the invention may have a thickness of from about 50 to about 500  $\mu m$ , preferably from about 75 to 300  $\mu m$ . Antioxidants, antistatic agents, plasticizers and other known additives may be incorporated into the support, if desired.

polyethylene-coated paper is employed.

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In order to improve the adhesion of the ink-receiving layer to the support, the surface of the support may be subjected to a corona-discharge treatment prior to applying the image-receiving layer.

Coating compositions employed in the invention may be applied by any number of well known techniques, including dip-coating, wound-wire rod coating, doctor blade coating, gravure and reverse-roll coating, slide coating, bead coating, extrusion coating, curtain coating and the like. Known coating and drying methods are described in further detail in Research Disclosure no. 308119, published Dec. 1989, pages 1007 to 1008. Slide coating is preferred, in which the base layers and overcoat may be simultaneously applied. After coating, the layers are generally dried by simple evaporation, which may be accelerated by known techniques such as convection heating.

To improve colorant fade, UV absorbers, radical quenchers or antioxidants may also be added to the image-receiving layer as is well known in the art. Other additives include pH modifiers, adhesion promoters, rheology modifiers, surfactants, biocides, lubricants, dyes, optical brighteners, matte agents, antistatic agents, etc. In order to obtain adequate coatability, additives known to those familiar with such art such as surfactants, defoamers, alcohol and the like may be used. A common level for coating aids is 0.01 to 0.30% active coating aid based on the total solution weight. These coating aids can be nonionic, anionic, cationic or amphoteric. Specific examples are described in MCCUTCHEON's Volume 1: Emulsifiers and Detergents, 1995, North American Edition.

The coating composition can be coated either from water or organic solvents, however water is preferred. The total solids content should be selected to yield a useful coating thickness in the most economical way, and for particulate coating formulations, solids contents from 10-40% are typical.

Ink jet inks used to image the recording elements of the present invention are well-known in the art. The ink compositions used in ink jet printing typically are liquid compositions comprising a solvent or carrier liquid, dyes or

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pigments, humectants, organic solvents, detergents, thickeners, preservatives, and the like. The solvent or carrier liquid can be solely water or can be water mixed with other water-miscible solvents such as polyhydric alcohols. Inks in which organic materials such as polyhydric alcohols are the predominant carrier or solvent liquid may also be used. Particularly useful are mixed solvents of water and polyhydric alcohols. The dyes used in such compositions are typically water-soluble direct or acid type dyes. Such liquid compositions have been described extensively in the prior art including, for example, U.S. Patents 4,381,946; 4,239,543 and 4,781,758, the disclosures of which are hereby incorporated by reference.

Although the recording elements disclosed herein have been referred to primarily as being useful for ink jet printers, they also can be used as recording media for pen plotter assemblies. Pen plotters operate by writing directly on the surface of a recording medium using a pen consisting of a bundle of capillary tubes in contact with an ink reservoir.

The following example is provided to illustrate the invention.

#### **EXAMPLE**

#### 20 Element 1 of the Invention

A coating solution was prepared by combining furned alumina (Cab-O-Sperse® PG003, Cabot Corp.), PVA-B and crosslinker 2,3-dihydroxy-1,4-dioxane (Clariant Corp.) in a ratio of 88:10:2 to give an aqueous coating formulation of 30% solids by weight, so that the crosslinking agent is present in an amount of 20 weight % of the poly(vinyl alcohol) binder.

The layer was bead-coated at  $40^{\circ}$ C on polyethylene-coated paper base which had been previously subjected to corona discharge treatment. The coating was then dried at  $60^{\circ}$ C by forced air to yield a recording element with a thickness of  $40 \ \mu m$ .

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## Element 2 of the Invention

This element was prepared the same as Element 1 except that the ratio of components was 87:10:3 to give an aqueous coating formulation of 30% solids by weight, so that the crosslinking agent is present in an amount of 30 weight % of the poly(vinyl alcohol) binder.

## Element 3 of the Invention

This element was prepared the same as Element 1 except that the ratio of components was 86:10:4 to give an aqueous coating formulation of 30% solids by weight, so that the crosslinking agent is present in an amount of 40 weight % of the poly(vinyl alcohol) binder.

#### Comparative Element C-1

This element was prepared the same as Element 1 except that

15 PVA-D was used instead of PVA-B, and the ratio of components was 84:15:1 to
give an aqueous coating formulation of 30% solids by weight, so that the
crosslinking agent is present in an amount of 6.6 weight % of the poly(vinyl
alcohol) binder.

## 20 Comparative Element C-2

This element was prepared the same as Element 1 except that PVA-D was used instead of PVA-B, and the ratio of components was 86.5:12.5:1 to give an aqueous coating formulation of 30% solids by weight, so that the crosslinking agent is present in an amount of 8 weight % of the poly(vinyl alcohol) binder.

#### Comparative Element C-3

This element was prepared the same as Element 1 except that PVA-D was used instead of PVA-B, and the ratio of components was 89:10:1 to give an aqueous coating formulation of 30% solids by weight, so that the

crosslinking agent is present in an amount of 10 weight % of the poly(vinyl alcohol) binder.

## Coating Quality

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The above dried coatings were visually evaluated for cracking with the following results:

Table 2

Recording Element	Cracking
1	None
2	None
3	None
C-1	None
C-2	None
C-3	Some

The above results show that neither any of the recording elements of the invention nor two comparative elements exhibited any cracking.

## Dry Time

Test images of cyan, magenta, yellow, red, green, blue and black bars, each 1.1 cm by 13.5 cm, were printed on the above elements using an Epson Stylus® Photo 870 using inks with catalogue number T008201. Immediately after ejection from the printer, a piece of bond paper was placed over the printed image and rolled with a smooth, heavy weight. Then the bond paper was separated from the printed image. Ink transferred to the bond paper if the recording element was not dry. The length of the bar imaged on the bond paper was measured and is proportional to the dry time. Dry times corresponding to a length of about 40 cm or less are acceptable.

Table 3

Recording Element	Proportional Dry Time
	(cm)
1	6
2	2
3	6
C-1	91
C-2	91
C-3	65

The above results show that the elements of the invention had much better dry times than all the comparative elements.

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## Coalescence

A test image of a green patch was printed on each of the above elements using an Epson Stylus® Photo 870 using inks with catalogue number T008201. Coalescence of the ink on the patches was visually rated as follows:

10 1 = None

2 = Slight

3 = Moderate

4 = Severe

The following results were obtained:

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Table 4

Recording Element	Coalescence
1	3
2	2
3	1
C-1	4
C-2	4
C-3	4

The above results show that the recording elements of the invention had much less coalescence than the comparative elements.

Although the invention has been described in detail with reference to certain preferred embodiments for the purpose of illustration, it is to be understood that variations and modifications can be made by those skilled in the art without departing from the spirit and scope of the invention.